DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Bulk Manufacturer of Controlled Substances Application: S&B Pharma Inc.

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before [INSERT DATE 60 DAYS AFTER DATE OF PUBLICATION IN THE FEDERAL REGISTER].

ADDRESS: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION:

The Attorney General has delegated his authority under the Controlled Substances

Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR

0.100(b). Authority to exercise all necessary functions with respect to the promulgation
and implementation of 21 CFR part 1301, incident to the registration of manufacturers,
distributors, dispensers, importers, and exporters of controlled substances (other than
final orders in connection with suspension, denial, or revocation of registration) has been
redelegated to the Assistant Administrator of the DEA Diversion Control Division

("Assistant Administrator") pursuant to section 7 of 28 CFR part 0, appendix to subpart

R.

In accordance with 21 CFR 1301.33(a), this is notice that on February 28, 2017, S&B Pharma Inc., 405 South Motor Avenue, Azusa, California 91702-3232 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled Substance	Drug Code	Schedule
Marihuana	7360	I
Tetrahydrocannabinols	7370	I
Amphetamine	1100	II
Methamphetamine	1105	II
Lisdexamfetamine	1205	II
Methylphenidate	1724	II
Pentobarbital	2270	II
4-Anilino-N-phenethyl-4-piperidine (ANPP)	8333	II
Tapentadol	9780	II
Fentanyl	9801	II

The company plans to manufacture the listed controlled substances in bulk for use in product development and for commercial sales to its customers. In reference to drug codes 7360 (marihuana) and 7370 (THC), the company plans to bulk manufacture both as synthetic substances. No other activity for these drug codes is authorized for this registration.

Dated: February 18, 2019.

John J. Martin,

Assistant Administrator.

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